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Extraction and Characterization of Cellulosic Nanowhisker obtained from Discarded Cotton Fibers

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Abstract

This study aimed to characterize cellulose nanowhiskers extracted from discarded cotton fibers. The nanocellulose was characterized by dynamic light scattering, zeta potential, Scanning Transmission Electron Microscopy (STEM), Atomic force microscopy (AFM) and X-ray powder diffraction (XRD). The nanocrystals are characterized by a rod-like morphology with sizes between 100 nm and 300 nm. The acidic hydrolyses applied to the extract nanocrystals increased the crystallinity index from 72 to 86%. The average length and diameter obtained were 98.1 ± 4.7 and 8.0 ± 3.4 nm, respectively. The aspect ratio of the fibers, around 12.3, was lower than the values usually reported in the literature. The dynamic light scattering results were in accordance with the nano crystal dimensions showing an average size of 235 nm. Cellulose nanowhiskers obtained from cotton rejected by textile industry due to their abundance, low cost, high crystallinity, strength, low weight and biodegradability, are the ideal candidates for the processing of polymer nanocomposites.

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1. Introduction

Cellulose is a homopolymer of β -1,4-d-glucose molecules linked in a linear chain that can be found in nature among various sources like plants, algae, marine creatures and bacteria. The single cotton fiber (thickness: 20 – 30 μm) consists of superfine fibrils having diameter in the range of nanometer.¹ Cellulose, in its native state (termed cellulose-I) in the cell walls of plants and trees, forms a crystalline structure, whereby chains aggregate via hydrogen bonding.¹ Two forms of crystalline cellulose are known to occur in plants namely cellulose Ia and Ib. All cellulose sources, however, are not completely crystalline. The presence of paracrystalline or amorphous cellulose is often significant, although it varies from species to species. The presence of this amorphous portion to cellulose makes the structure susceptible to acid hydrolysis, and the eventual breakdown into individual crystallites.² Cellulose nanocrystals (CNC), sometimes referred as nanofibers, cellulose crystallites or simply crystals, are the crystalline domains of cellulose sources, with structure in nanoscale dimensions.³ Nanowhiskers are defined as the nanocellulose with diameters ranging from 2 nm to 20 nm and length ranging from 100 nm to 2.1 μm .⁴ These nanomaterials can be isolated from several natural fibers and are characterized by their low cost, abundance and biodegradability.⁵ When isolated, they can be applied in a wide range of applications in different fields, such as, engineered nanocomposites, construction materials, porous materials, intelligent or active package, functional surface among others.⁶ According to literature the main properties of nanocellulose materials are: biodegradability, biocompatibility, high strength, high surface area, moisture absorption, dimensional stability and chemical functionality.^{7,8} Cotton is one of the natural fibers with highest percentage of cellulose (>95%) and it was widely used as source of nanocellulose.⁹ However, the information about cotton nanowhiskers production methodologies and physico-chemical properties are scarce.¹⁰⁻¹² Thus, the aim of this work was to extract and characterize the cellulose nanowhisker obtained from cotton rejected by textile industry. The physical and chemical properties of CNC obtained were studied by analysis of surface charge, crystal size dimension and morphology, crystallinity and chemical structure.

2. Materials and methods

2.1. Cellulose nanowhiskers preparation

A discarded 100% cotton fabric (textile industry residue) was used in this study. Cotton residues were milled and hydrolyzed by alkali treatment in a solution of 5% of NaOH at 80 °C. Then a bleaching process was applied in a bath composed by 3mL/L of H_2O_2 200V, 2mL/L of NaOH 38°Be, 1mL/L of stabilizer and 1mL/L of detergent. Finally, the method of acidic hydrolysis was applied with minor adaptations using a solution of 60% of H_2SO_4 . At the end of process, the reaction was stopped by adding 10 times the volume of cold deionized water. The colloidal suspension was centrifuged for 20 minutes at 13.000 rpm in a High-speed Refrigerated Centrifuge. The suspension obtained was ultra-sonicated to prevent agglomerates formation and dialyzed with tap water during 24 h until pH of 6-7 was reached. The process from centrifugation through dialysis was repeated three times.

2.2. Dynamic Light Scattering (DLS)

The particle size distribution (z-average-size) and the polydispersity index (PDI) of the silver nanoparticles were determined with a ZetaPlus instrument (Brookhaven Instruments Corporation). The samples were measured at a constant temperature of 25 ± 1 °C. The results were averaged over 30 runs. Each value was obtained by averaging measurements of three samples.

2.3. Zeta Potential Measurements

The z-potentials were determined in an electrophoretic lightscattering ZetaPlus apparatus (Brookhaven Instruments Corporation) to examine nanocellulose colloidal stability and charge. The electrophoretic determinations of z-potentials were made in aqueous media at moderate electrolyte concentration. Each value was obtained by averaging measurements of three samples.

2.4. Scanning Transmission Electron Microscopy (STEM)

STEM analysis of the nanowhiskers samples were carried out with an Ultra-high resolution Field Emission Gun Scanning Electron Microscopy (FEG-SEM), NOVA 200 Nano SEM, FEI Company. Secondary electron images were performed with an acceleration voltage between 5 and 10 kV. Backscattering Electron Images were realized with an acceleration voltage of 15 kV. The synthesized nanowhiskers suspension was dropped in copper grids with carbon film 400 meshes, 3mm diameter. The average size was extrapolated measuring the average diameter based in 50 observations.

2.5. X-ray powder diffraction (XRD)

The crystalline phases of cellulose nanowhiskers were determined by using of X-ray diffractometer Shimadzu model: XRD 6000 with CuK α radiation in the range 20-70° ($\lambda=0.154\text{nm}$).

2.6. Atomic force microscopy (AFM)

AFM experiments were performed in a tapping mode in air using a Multimode AFM from Agilent, model 5500, at room temperature. Silicon cantilevers (AppNano) with a constant force between 25 and 75 N m⁻¹ and a resonance frequency of 339 kHz were used. The PicoView 1.10 software version was used for the simultaneous recording of the topography, phase and amplitude magnitudes of the images.

3. Results and discussion

The amorphous regions of the cotton structure are the main target for the cleavage of the microfibrils into short monocrystals under acid hydrolysis due to their faster hydrolysis kinetics. The used method revealed to be a useful procedure for the preparation of highly crystalline particles of nanowhiskers. Under the used controlled conditions it was possible an efficient disruption of the amorphous regions surrounding and embedded within cellulose microfibrils leaving the microcrystalline segments intact. Since the geometrical characteristics of cellulose whiskers depend on the origin of cellulose microfibrils and acid hydrolysis process conditions such as time, temperature, and purity of materials, the morphological and physico-chemical characterization of nanowhiskers is extremely important for its potential uses as nanocomposite.^{13,14}

The results of the zeta potential show that the nanocrystals are quite stables displaying an absolute value of 25.35±1.5 mV (Fig. 1-a). A value of 25 mV is widely reported in literature and it is considered the minimum value for stable nanocellulose dispersions.¹⁵ The particle size distribution of the nanosizer analysis is remarkably monodisperse showing a single sharp peak in intensity with sizes between 100 and 300 nm and an average size of 235 nm (Fig. 1-b). However, in the case of rod-like structure the dynamic light scattering display some limitations on the size estimation (Fig. 1-c). A more sensitive methodology like AFM will be necessary to confirm this data. The formation of a precipitate of insoluble clusters of nanocellulose particles was also observed. The analysis of the precipitate exhibits a wide peak indicating a high dispersion of sizes in micro range.

Fig.1. Nanosizer and Zeta potential results.

The XRD patterns demonstrated the high crystallinity of the extracted cellulose nanowhiskers. The diffractograms of cellulose and nanowhiskers show the characteristic peaks of cellulose I at 22.5° (200) and 34.5° (040).¹⁶ The superimposed lattice peaks between 14.8° (110) and 16.7° (110) are better defined in the nanowhiskers cellulose than cellulose fibers. The higher crystallinity was confirmed by the calculation of the crystallinity index of the nanowhiskers (86%) compared to the cellulose (72%). An increase of 14% of crystallinity is in accordance with the available literature about the use of nanowhiskers as nanofiller for composites.^{9,17,18}

Fig 2. XRD measures of cellulose and nanowhiskers

The average length and diameter were estimated by AFM and STEM analysis with values of 98.1 ± 4.7 and 8.0 ± 3.4 nm, respectively (Fig. 3). The aspect ratio of the fibers, around 12.3, was lower than the values usually reported in literature.^{19,20} TEM picture reveals that cellulose nanowhiskers have a high surface area to volume ratio. This means that the surface could play a dominant role in the mechanical efficiency of stress transfer in a nanocomposite as well as in the ability to modify the surface chemistry. STEM picture reveal zone of agglomeration of nanocellulose with dispersed crystallites and individual crystals.

Fig.3. AFM topography images (2x2 μm) and STEM of the nanowhiskers.

Conclusion

Cellulose nanowhiskers were successfully extracted by acid hydrolysis from cotton residues showing an aspect ratio of 12 and high crystallinity of 86%. Nanocrystals sizes and dispersion was accurately measured by TEM, STEM and light scattering analysis. The results presented in this work suggest that the acidic route is an efficient alternative for obtaining monodisperse cellulose nanowhiskers from industrial cotton wastes.

References

1. Satyamurthy, P., et al., *Carbohydrate Polymers* (2011) **83** (1), 122
2. Eichhorn, S. J., *Soft Matter* (2011) **7** (2), 303
3. Azizi Samir, M. A. S., et al., *Biomacromolecules* (2005) **6** (2), 612
4. Rosa, M. F., et al., *Carbohydrate Polymers* (2010) **81** (1), 83
5. Eichhorn, S. J., *Soft Matter* (2011) **7** (2), 303
6. Siró, I., and Plackett, D., *Cellulose* (2010) **17** (3), 459
7. Hossain, K. M. Z., et al., *Soft Matter* (2012) **8** (48), 12099
8. Siqueira, G., et al., *Biomacromolecules* (2009) **10** (2), 425
9. Morais, J. P. S., et al., *Carbohydrate Polymers* (2013) **91** (1), 229
10. Adel, A. M., et al., *Bioresource Technol* (2010) **101** (12), 4446
11. Adel, A. M., et al., *Carbohydrate Polymers* (2011) **83** (2), 676
12. Gardner, D. J., et al., *J Adhes Sci Technol* (2008) **22** (5-6), 545
13. Bras, J., et al., *Carbohydrate Polymers* (2011) **84** (1), 211
14. Cao, X. W., et al., *Carbohydrate Polymers* (2012) **90** (2), 1075
15. Olszewska, A., et al., *Soft Matter* (2013) **9** (31), 7448
16. Elazzouzi-Hafraoui, S., et al., *Biomacromolecules* (2008) **9** (1), 57
17. Cherian, B. M., et al., *Journal of Agricultural and Food Chemistry* (2008) **56** (14), 5617
18. de Morais Teixeira, E., et al., *Cellulose* (2010) **17** (3), 595
19. Abdul Khalil, H. P. S., et al., *Carbohydrate Polymers* (2012) **87** (2), 963
20. Lavoine, N., et al., *Carbohydrate Polymers* (2012) **90** (2), 735